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(54) Mercury vapour separation

(54) Mercury vapor in a gas can easily be removed from the gas by contacting the gas with an adsorbent comprising an activated carbon having as supported thereon (1) one or more components included in one of three groups shown below and (2) one or more components included in the remaining two groups shown below.

Group	Component
I	sulfur
II	sulfate and nitrate of Al, V, Fe, Co, Ni, Cu, Zn or NH ₄
III	oxide of iodine, oxyacid corresponding to the oxide of iodine, salt of said oxyacid, and bromide and iodide of K, Na or NH ₄

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SPECIFICATION

Process for removal of mercury vapor and adsorbent therefor

The present invention relates to a process for the removal of mercury vapor and an adsorbent thereof.

5 Electrolytic hydrogen is used in process for manufacturing various products such as pharmaceutical and food products, whereby trace amounts of mercury vapor act as catalyst poison and also contaminate such products, thus constituting in some instances the cause of much argument. Mercury vapor contained in natural gas corrodes heat exchanger made of aluminum and other types of equipment in the process of gas liquefaction, and is responsible for serious disasters. Also, mercury vapor contained in incinerator waste gases and waste gases from factories handling mercury is regarded as controversial from the standpoints of air pollution control and working environment preservation. 10

Therefore, it becomes necessary to remove mercury vapor in these gases by means of some appropriate treatment. As the method of removing mercury vapor in gas, there have been conventionally known the process of washing with chemical solutions and the method of dry adsorbing with use of adsorbents such as activated carbon and ion exchange resins. Nevertheless, the first-mentioned process of washing with chemical solutions suffers from the defects that the efficiency of removal of mercury vapor is poor and that waste solutions are produced, while the last-mentioned dry adsorption method, with its respectively small capacity to adsorb mercury vapor, is far from satisfactory. 15

20 The present inventors, in view of such circumstances, conducted extensive research, and as a result, have come to the finding that the adsorbents, which consist of activated carbon having as supported thereon two or more specific components selected from (I) sulfur, (II) specific salt components and (III) specific halogen compounds, adsorb efficiently mercury vapor in gases and also exhibit an outstandingly increased adsorption capacity. 20

25 Thus the present invention relates to:

A process for the removal of mercury vapor from a gas containing the same, which comprises contacting the gas with an adsorbent comprising activated carbon having as supported thereon (1) one or more components included in one of the three groups shown below and (2) one or more components included in the remaining two groups shown below:

Group	Component
I	sulfur
II	sulfate and nitrate of Al, V, Fe, Co, Ni, Cu, Zn or NH ₄
III	oxide of iodine, oxyacid corresponding to the oxide of iodine, salt of said oxyacid, and bromide and iodide of K, Na or NH ₄

and an adsorbent used in the above process.

The adsorbent of the present invention can be prepared by having (1) one or more components included in one of the three groups shown above and (2) one or more components included in the remaining two groups shown above supported on activated carbon.

35 The activated carbon, which is used in this case, may be any type, so long as in manufactured by a known process from wood charcoal, coal, coke, coconut shells, resins, etc. employed as raw materials and exhibits a specific surface area of 200 to 2000 m²/g. 35

Examples of the components of Group II include sulfates such as (NH₄)₂SO₄, Al₂(SO₄)₃, VOSO₄, FeSO₄, Fe₂(SO₄)₃, CoSO₄, NiSO₄, CuSO₄, ZnSO₄, etc. and nitrates such as NH₄NO₃, Al(NO₃)₃, VO(NO₃)₂, Fe(NO₃)₂, Fe(NO₃)₃, CO(NO₃)₂, Ni(NO₃)₂, Cu(NO₃)₂, Zn(NO₃)₂, etc. 40

Examples of the components of Group III include oxides of iodine such as I₂O₅, IO₂, etc., oxyacids corresponding to the oxide of iodine such as HIO₂, HIO₃, etc., salts of said oxyacids such as alkali metal salts (e.g. NaIO₂, KIO₂, NaIO₃, KIO₃, NaIO₄, KIO₄), alkaline earth metal salts (e.g. Mg(IO₃)₂, Ca(IO₃)₂, Sr(IO₃)₂), ammonium salts (e.g. NH₄IO₃, NH₄IO₄), etc., bromide such as KBr, NaBr, NH₄Br, etc., and iodide such as KI, NaI, NH₄I, KI₃, etc. 45

When sulfur of Group I is used, the amount of sulfur to be supported is 10 to 800 mg per g of activated carbon, preferably 20 to 600 mg. When the component or components of Group II is used, the amount of component or total amount of components of Group II is 5 to 500 mg per g of activated carbon, preferably 10 to 400 mg. When the component or components of Group III is used, the amount

of the component or total amount of components of Group III is 1 to 500 mg per g of activated carbon, preferably 2 to 400 mg, most preferably 10 to 400 mg.

In order to allow activated carbon to support the components selected from Groups I, II and III, in the manufacture of the adsorbent for mercury vapor according to the present invention, there may be mentioned, by way of example, the method which comprises dissolving or suspending these components to be supported together in water or a water-soluble solvent and impregnating or spraying activated carbon with the resultant solution or suspension, followed by heating at 110 to 400°C, if necessary. Also, each of these components may be supported on activated carbon, individually, and the order thereof may be arbitrary.

The heating procedure as described above can be carried out for example in the presence of nitrogen gas, carbon dioxide gas, steam, air, oxygen-containing gas, combustion gas or gas mixtures thereof. As the means of heating, any procedure may be employable, and use is desirably made for example of the method which comprises contacting the above-mentioned gas with activated carbon under heating. As the mode of contacting employable in this case, there may be mentioned the fixed-bed, fluidized bed, slurry and rotary-kiln types and others. The heating time varies with the kind and temperature of the surrounding gases, etc., and is normally not less than 10 minutes, preferably 1/2 to 24 hours.

With reference to the adsorbents for mercury vapor according to the present invention, the adsorbent having component of Group I, component(s) included in Group II and component(s) included in Group III exhibits greater removal efficiency and adsorption capacity for mercury than the one having component(s) included in one of the three groups and component(s) included in one of the remaining two groups.

Further, the adsorbent of the present invention may be an activated carbon having as supported thereon one or more of sulfur compounds in addition to the components included in Group I, II and III. Examples of the sulfur compound include thiocyanates such as NH_4SCN , NaSCN , KSCN , $\text{Mg}(\text{SCN})_2$, $\text{Ca}(\text{SCN})_2$ and $\text{Sr}(\text{SCN})_2$, thiosulfate such as $(\text{NH}_4)_2\text{S}_2\text{O}_3$, $\text{Na}_2\text{S}_2\text{O}_3$, $\text{K}_2\text{S}_2\text{O}_3$, MgS_2O_3 , CaS_2O_3 and SrS_2O_3 , sulfamates such as $\text{NH}_4\text{OSO}_2\text{NH}_2$, $\text{NaOSO}_2\text{NH}_2$, KOSO_2NH_2 , $\text{Mg}(\text{OSO}_2\text{NH}_2)_2$, $\text{Ca}(\text{OSO}_2\text{NH}_2)_2$ and $\text{Sr}(\text{OSO}_2\text{NH}_2)_2$, and sulfides such as $(\text{NH}_4)_2\text{S}$, Na_2S , K_2S , MgS , CaS and SrS . The total amount of the sulfur compounds is usually 10 to 800 mg per g of activated carbon, preferably 20 to 600 mg.

The process for the removal of mercury vapor of the present invention is carried out by contacting a gas containing mercury vapor with the adsorbent of the present invention. The contacting temperature is not more than 150°C, preferably -10 to 120°C, more preferably 0 to 120°C, and the contacting pressure is not more than 100 mg/cm², preferably 0.1 to 70 kg/cm², while the contacting time at 25°C and at 1 kg/cm² is 1/10 to 30 seconds, preferably 1/5 to 20 seconds. Also, contact between the said adsorbent and the gas containing vaporized mercury can be effected for example by means of the fixed-bed, moving-bed and fluidized-bed operations.

Further, the present inventors have found that an activated carbon having as supported thereon sulfur is desirably prepared by mixing activated carbon with finely powdered sulfur and heating the resultant mixture at 110 to 350°C, preferably 110 to 300°C.

The fine particles of sulfur, which is used in this preparation, may be any of fine pulverized sulfur, if it shows a grain size of not more than 8 mesh, preferably not more than 10 mesh. The amount of sulfur to be supported onto activated carbon is normally 10 to 800 mg per g of activated carbon. In the preparation process, the first step is conducted into practice by mixing activated carbon uniformly with fine particles of sulfur. Such procedure of mixing may be carried out by mixing directly activated carbon with fine particles of sulfur, whereby water or solvent mixtures of water-soluble solvents (e.g., alcohols, etc.) with water may be desirably employed. In the case of the use of water or solvent, examples of the employable procedure include a procedure which comprises firstly suspending fine particles of sulfur in water or solvent and impregnating or spraying activated carbon with the resultant suspension and a procedure which comprises impregnating activated carbon with water or solvent and coating it with fine particles of sulfur. In case of preparing an activated carbon having as supported thereon sulfur and other component or components defined in the present invention, it is preferable to dissolve or suspend in advance in water or solvent as mentioned above the components intended to be supported onto activated carbon. Each amount of the components to be supported is as defined above. The means of mixing may be any of means, such as mixer, sugar-coating machine and agitator, which may be utilized for mixing solids. A mixture of activated carbon and fine particles of sulfur, along with other components, if necessary, is subsequently heated at 110°C to 350°C, preferably 110°C to 300°C. Heating can be effected for example in the presence of nitrogen gas, carbon dioxide gas, steam, combustion gas, air, oxygen-containing gas or gas mixtures thereof. The means of heating may be any of means, whereby, for example, the procedure comprising heating the above gases and contacting it with a mixture of activated carbon and sulfur is desirably employed. As the mode or type of contacting employable in this case, there may be mentioned fixed-bed, moving-bed, fluidized-bed, slurry and rotary-kiln types, and other. The heating time varies with the kinds and temperatures of the surrounding gases, etc., and is normally not less than 30 minutes, preferably 1 to 240 hours, more preferably 1 to 50 hrs. It should be noted that when the above heating temperature is lower than 110°C, sulfur fails to reach the inner surface of activated carbon and therefore, the activated carbon having as supported

thereon sulfur showing excellent adsorption effect cannot be obtained. Conversely, the increased heating temperature in excess of 350°C causes sulfur vapor to be generated, and is consequently unfavorable. The activated carbon having as supported thereon sulfur as obtained by the above procedure has sulfur supported uniformly on the pore surface of activated carbon, and possesses excellent adsorption capacity for mercury vapor and so forth.

EXAMPLE 1

Activated carbon A₀ of 16 to 24 mesh having the BET specific surface area of 1230 m²/g was evenly sprayed with an aqueous solution or suspension of the given amount of the components, other than sulfur, shown in the following table. When sulfur is used, the given amount of finely powdered sulfur was mixed with the activated carbon. The activated carbon was then heated in an air at 110°C.

(Adsorbents (1)—(64))

As controls, activated carbon A₀ was mixed, in the same procedure as described above, with the given amount each of the components alone, and heated at 110°C. (Adsorbents A₁ through A₁₆)

Six ml each of the Adsorbents A₀ through A₁₆ and (1) through (64) as obtained by the above procedures were packed into a 1.6 cm ϕ glass column, respectively, and nitrogen gas of 25°C containing 9 mg/m³ of mercury vapor was passed through each column at a linear flow rate of 40 cm/sec to conduct the breakthrough adsorption tests for mercury vapor. The results are as shown in Table 1.

TABLE 1-1

Adsorbent No.	The first component or additive; and its amount supported (mg/g)	The second component; and its amount supported (mg/g)	The third component; and its amount supported (mg/g)	Percent removal of mercury vapor at the elapsed time shown below (%)			
				5 hrs.	10 hrs.	20 hrs.	50 hrs.
A ₀ (Control)	0	0	0	0	0	0	0
A ₁ (")	0	0	NH ₄ I - 150	53	28	3	0
A ₂ (")	0	0	NaI - 150	59	35	3	0
A ₃ (")	0	0	KI - 150	78	51	15	0
A ₄ (")	0	0	KBr - 150	48	35	7	0
A ₅ (")	0	Al ₂ (SO ₄) ₃ - 100	0	21	3	0	0
A ₆ (")	0	FeSO ₄ - 100	0	66	42	28	7
A ₇ (")	0	CuSO ₄ - 100	0	89	61	29	8
A ₈ (")	0	CoSO ₄ - 100	0	55	40	23	10
A ₉ (")	0	NH ₄ NO ₃ - 100	0	33	9	0	0
A ₁₀ (")	0	Fe(NO ₃) ₃ - 100	0	96	84	50	11
A ₁₁ (")	0	(NH ₄) ₂ SO ₄ - 100	0	19	5	0	0
A ₁₂ (")	S - 100	0	0	68	50	38	24
A ₁₃ (")	S - 200	0	0	51	41	30	25
A ₁₄ (")	0	ZnSO ₄ - 100	0	35	5	0	0
A ₁₅ (")	0	Zn(NO ₃) ₂ - 100	0	88	74	43	10
A ₁₆ (")	0	0	HI - 100	85	13	0	0

TABLE 1-2

Adsorbent No.	The first component or additive; and its amount supported (mg/g)	The second component; and its amount supported (mg/g)	The third component; and its amount supported (mg/g)	Percent removal of mercury vapor at the elapsed time shown below (%)			
				5 hrs.	10 hrs.	20 hrs.	50 hrs.
(1) This (invention)	S - 100	0	NH ₄ I - 10	93	85	59	30
(2) (")	S - 100	0	NaI - 10	90	72	48	31
(3) (")	S - 100	0	KI - 10	94	78	45	29
(4) (")	S - 100	0	KBr - 10	91	71	50	42
(5) (")	S - 100	0	NH ₄ Br - 10	88	73	49	41
(6) (")	S - 100	0	KI - 5 KBr - 5	94	85	61	49
(7) (")	S - 100	0	NaI - 5 KBr - 5	92	73	47	41
(8) (")	S - 100	0	KI - 2	81	65	41	31
(9) (")	S - 150	0	KI - 5	83	66	40	33
(10) (")	S - 200	0	KI - 5	80	67	43	35
(11) (")	S - 100	Al(SO ₄) ₃ - 10	0	88	71	45	32
(12) (")	S - 100	FeSO ₄ - 10	0	95	78	51	23
(13) (")	S - 100	CuSO ₄ - 10	0	95	88	69	48

TABLE 1-3

Adsorbent No.	The first component or additive; and its amount supported (mg/g)	The second component; and its amount supported (mg/g)	The third component; and its amount supported (mg/g)	Percent removal of mercury vapor at the elapsed time shown below (%)			
				5 hrs.	10 hrs.	20 hrs.	50 hrs.
(14) This (invention)	S - 100	CoSO ₄ - 10	0	89	75	61	38
(15) (")	S - 100	NiSO ₄ - 10	0	90	75	63	41
(16) (")	S - 100	VO SO ₄ - 10	0	99	70	48	39
(17) (")	S - 100	(NH ₄) ₂ SO ₄ - 10	0	81	61	51	39
(18) (")	S - 100	Fe(NO ₃) ₂ - 10	0	93	67	63	44
(19) (")	S - 100	Co(NO ₃) ₂ - 10	0	96	75	64	40
(20) (")	S - 100	FeSO ₄ - 5	0	89	75	60	33
(21) (")	S - 150	FeSO ₄ - 5	0	88	76	63	38
(22) (")	S - 200	FeSO ₄ - 5	0	90	78	65	41
(23) (")	S - 100	ZnSO ₄ - 10	0	80	71	43	37
(24) (")	S - 100	FeSO ₄ - 5 CuSO ₄ - 5	0	96	80	52	41
(25) (")	S - 100	FeSO ₄ - 5 Fe(NO ₃) ₃ - 5	0	91	73	60	43

TABLE 1-4

Adsorbent No.	The first component or additive; and its amount supported (mg/g)	The second component; and its amount supported (mg/g)	The third component; and its amount supported (mg/g)	Percent removal of mercury vapor at the elapsed time shown below (%)			
				5 hrs.	10 hrs.	20 hrs.	50 hrs.
(26) (This) invention	0	(NH ₄) ₂ SO ₄ - 50	I ₂ O ₅ - 50	100	100	100	85
(27) (..)	0	Al ₂ (SO ₄) ₃ - 50	I ₂ O ₅ - 50	100	100	100	98
(28) (..)	0	Fe ₂ (SO ₄) ₃ - 50	I ₂ O ₅ - 50	100	100	100	93
(29) (..)	0	Fe(NO ₃) ₃ - 50	I ₂ O ₅ - 50	100	100	100	96
(30) (..)	0	Co(NO ₃) ₂ - 50	I ₂ O ₅ - 50	100	100	100	97
(31) (..)	0	Ni(NO ₃) ₂ - 50	HIO ₃ - 50	100	100	100	98
(32) (..)	S - 50	0	I ₂ O ₅ - 50	100	100	100	95
(33) (..)	S - 50	VOSO ₄ - 50	I ₂ O ₅ - 50	100	100	100	100
(34) (..)	S - 50	FeSO ₄ - 50	I ₂ O ₅ - 50	100	100	100	100
(35) (..)	NaOSO ₂ NH ₂ - 50	ZnSO ₄ - 50	I ₂ O ₅ - 50	100	100	100	100
(36) (..)	K ₂ S ₂ O ₃ - 50	Cu(NO ₃) ₂ - 50	HIO ₃ - 50	100	100	100	100
(37) (..)	K ₂ S - 50	FeSO ₄ - 50	I ₂ O ₅ - 50	100	100	100	100
(38) (..)	0	FeSO ₄ - 50	KIO ₄ - 50	100	100	100	63
(39) (..)	S - 50	FeSO ₄ - 50	Mg(IO ₃) ₂ - 50	100	100	100	100

TABLE 1-4

Adsorbent No.	The first component or additive; and its amount supported (mg/g)	The second component; and its amount supported (mg/g)	The third component; and its amount supported (mg/g)	Percent removal of mercury vapor at the elapsed time shown below (%)			
				5 hrs.	10 hrs.	20 hrs.	50 hrs.
(40) This (invention)	0	(NH ₄) ₂ SO ₄ - 50	NH ₄ I - 75	100	100	100	75
(41) (")	0	Al ₂ (SO ₄) ₃ - 50	NaI - 75	100	100	100	88
(42) (")	0	VOSO ₄ - 50	KI - 75	100	100	100	95
(43) (")	0	CoSO ₄ - 50	KBr - 50	100	100	100	90
(44) (")	0	FeSO ₄ - 50	NH ₄ Br - 75	100	100	100	88
(45) (")	0	FeSO ₄ - 50	KI - 75	100	100	100	93
(46) (")	0	CoSO ₄ - 50	NH ₄ I - 75	100	100	100	89
(47) (")	0	NiSO ₄ - 50	KI - 75	100	100	100	88
(48) (")	0	CuSO ₄ - 50	KIO ₃ - 75	100	100	100	83
(49) (")	0	ZnSO ₄ - 50	NaBrO ₃ - 75	100	100	100	80
(50) (")	0	CuSO ₄ - 50	NH ₄ Br - 75	100	100	100	95
(51) (")	0	FeSO ₄ - 50	NaI - 75	100	100	100	78
(52) (")	0	NH ₄ NO ₃ - 50	KI - 75	100	100	100	73
(53) (")	0	Al(NO ₃) ₃ - 50	NH ₄ I - 75	100	100	100	86
(54) (")	0	Fe(NO ₃) ₃ - 50	NaI - 75	100	100	100	81

TABLE 1-6

Adsorbent No.	The first component or additive; and its amount supported (mg/g)	The second component; and its amount supported (mg/g)	The third component; and its amount supported (mg/g)	Percent removal of mercury vapor at the elapsed time shown below (%)			
				5 hrs.	10 hrs.	20 hrs.	50 hrs.
(55) (This invention)	0	$\text{Fe}(\text{NO}_3)_3$ - 50	KBr - 75	100	100	100	83
(56) (")	0	$\text{Fe}(\text{NO}_3)_3$ - 50	NaBr - 75	100	100	100	82
(57) (")	0	$\text{Co}(\text{NO}_3)_2$ - 50	NH_4I - 75	100	100	100	91
(58) (")	0	$\text{Ni}(\text{NO}_3)_2$ - 50	NaI - 75	100	100	100	89
(59) (")	0	$\text{Cu}(\text{NO}_3)_2$ - 50	KI - 75	100	100	100	96
(60) (")	0	$\text{Zn}(\text{NO}_3)_2$ - 50	KI - 75	100	100	100	91
(61) (")	0	FeSO_4 - 100	KI - 400	100	100	100	100
(62) (")	0	FeSO_4 - 100	KI - 200	100	100	100	100
(63) (")	0	FeSO_4 - 100	KI - 50	100	100	100	65
(64) (")	0	FeSO_4 - 100	KI - 10	100	100	100	8

EXAMPLE 2

A 6 ml portion each of the Adsorbents A_1 , A_3 , A_5 , A_6 , A_9 , A_{10} , A_{14} , (28), (31), (32), (34), (36), (40), (42), (45), (48), (52), (54), (56), and (59) of Example 1 was packed into a 1.6 cm ϕ glass column, respectively, and hydrogen gas of 25°C containing 1 mg/m³ of Mercury vapor was passed through each column at a linear flow rate of 40 cm/sec to conduct the breakthrough adsorption tests for mercury vapor. The result are shown in Table 2.

5

TABLE 2

Adsorbent No.	Percent removal of mercury vapor at the elapsed time shown below (%)			
	50 hrs.	100 hrs.	150 hrs.	200 hrs.
A_1 (control)	42	15	6	0
A_3 (..)	65	35	11	3
A_5 (..)	42	32	8	0
A_6 (..)	58	40	13	9
A_9 (..)	26	15	0	0
A_{10} (..)	78	43	15	0
A_{14} (..)	51	23	1	0
(28) (This invention)	100	100	100	98
(31) (..)	100	100	100	100
(32) (..)	100	100	100	100
(34) (..)	100	100	100	100
(36) (..)	100	100	100	100
(40) (..)	100	100	100	75
(42) (..)	100	100	100	100
(45) (..)	100	100	100	100
(48) (..)	100	100	100	100
(52) (..)	100	100	100	83
(54) (..)	100	100	100	91
(56) (..)	100	100	100	95
(59) (..)	100	100	100	100

EXAMPLE 3

A 6 ml portion each of the Adsorbents A_1 , A_3 , A_5 , A_6 , A_{10} , A_{14} , (28), (31), (32), (34), (36), (40), (42), (45), (48), (52), (54), (56) and (59) of Example 1 was packed into a 1.6 cm ϕ glass column, respectively, and a gas (N—84.8 vol.%, O₂—14.5 vol.%, H₂O—0.7 vol.%) of 25°C containing 9 mg/m³ of mercury vapor was passed through each column at a linear flow rate of 40 cm/sec to conduct the breakthrough adsorption tests for mercury vapor. The results are as shown in Table 3.

10

10

TABLE 3

Adsorbent No.	Percent removal of mercury vapor at the elapsed time shown below (%)			
	50 hrs.	100 hrs.	150 hrs.	200 hrs.
A ₁ (Control)	68	39	18	8
A ₃ (..)	73	40	24	12
A ₅ (..)	63	39	18	10
A ₆ (..)	63	51	21	15
A ₁₀ (..)	32	18	4	0
A ₁₄ (..)	60	35	8	3
(28) (This invention)	100	100	100	100
(31) (..)	100	100	100	100
(32) (..)	100	100	100	100
(34) (..)	100	100	100	100
(36) (..)	100	100	100	100
(40) (..)	100	100	100	88
(42) (..)	100	100	100	100
(45) (..)	100	100	100	100
(48) (..)	100	100	100	100
(52) (..)	100	100	100	100
(53) (..)	100	100	100	100
(56) (..)	100	100	100	100
(59) (..)	100	100	100	100

EXAMPLE 4

5 A 6 ml portion each of the Adsorbents A₃, A₆, A₁₂, (2) through (4) and (12) through (14) of Example 1 was packed into a 1.6 cm³ glass column, and methane gas of 70°C containing 9 mg/m³ of mercury vapor was passed through each column at a linear flow rate of 40 cm/sec to conduct the breakthrough adsorption tests for mercury vapor at 70°C. The results are as shown in Table 4.

TABLE 4

Adsorbent No.	Percent removal of mercury vapor at the elapsed time shown below (%)						
	5 hrs.	10 hrs.	20 hrs.	50 hrs.	100 hrs.	150 hrs.	200 hrs.
A ₃ (Control)	58	59	63	63	35	30	23
A ₆ (")	20	38	40	40	39	37	37
A ₁₂ (")	55	58	65	68	68	70	70
(2) (This invention)	95	95	96	96	97	96	96
(3) (")	96	98	98	98	98	98	99
(4) (")	95	96	96	96	96	97	97
(12) (")	97	98	98	99	99	100	100
(13) (")	98	99	100	100	100	100	100
(14) (")	95	97	98	98	98	98	98

EXAMPLE 5

Activated carbon a₀ of 16 to 24 mesh having the BET specific surface area of 1050 m²/g was sprayed with a solution containing respectively given amount of sulfur (first component), a sulfate or nitrate (second component) and bromide or iodide (third component) (in the case of sulfur, a suspension was used), and subsequently dried at 120°C (Adsorbents (65) through (80)).

Six ml each of the Adsorbents a₀ and (65) through (80) was packed into a 1.6 cm³ glass column, respectively, and nitrogen gas of 25°C containing 9 mg/m³ of mercury vapor was passed through each column at a linear flow rate of 40 cm/sec to conduct the breakthrough adsorption tests for mercury vapor. The results are shown in Table 5.

TABLE 5

Adsorbent No.	The first component or additive; and its amount supported (mg/g)	The second component; and its amount supported (mg/g)	The third component; and its amount supported (mg/g)	Percent removal of mercury vapor at the elapsed time shown below (%)			
				5 hrs.	10 hrs.	20 hrs	50 hrs.
a ₀ (Control)	0	0	0	0	0	0	0
(65) (This invention)	S — 150	0	KI — 100	91	70	40	16
(66) (")	S — 150	FeSO ₄ — 100	0	89	65	40	11
(67) (")	S — 150	Zn(NO ₃) ₂ — 100	0	68	43	21	7
(68) (")	S — 150	0	NH ₄ I — 100	90	81	54	15
(69) (")	S — 75	FeSO ₄ — 50	KBr — 75	100	100	100	100
(70) (")	S — 75	FeSO ₄ — 50	NaI — 75	100	100	100	100
(71) (")	S — 75	FeSO ₄ — 50	KI — 75	100	100	100	100
(72) (")	S — 75	CuSO ₄ — 50	KI — 75	100	100	100	100
(73) (")	S — 75	Zn(NO ₃) ₂ — 50	KI — 75	100	100	100	100
(74) (")	S — 50	VOSO ₄ — 50	NH ₄ I — 75	100	100	100	100
(75) (")	S — 50	Fe(NO ₃) ₃ — 50	KIO ₃ — 75	100	100	100	100
(76) (")	S — 50	(NH ₄) ₂ SO ₄ — 50	NH ₄ Br — 75	100	100	100	100
(77) (")	S — 400	FeSO ₄ — 50	KI — 50	100	100	100	100
(78) (")	S — 100	FeSO ₄ — 50	KI — 50	100	100	100	100
(79) (")	S — 50	FeSO ₄ — 50	KI — 50	100	100	100	100
(80) (")	S — 10	FeSO ₄ — 50	KI — 50	100	100	100	99

EXAMPLE 6

From activated carbon A of 16 to 24 mesh having the BET surface area of 1100 m²/g, there was prepared the activated carbon having as supported thereon 100 mg/g of sulfur by the following procedure:

- 5 (1) Activated carbon having as supported thereon sulfur B₀ through B₃: 5
 10.0 g of finely powdered sulfur of not more than 50 mesh was uniformly suspended in 90 ml of water, and the suspension was sprayed evenly onto 100 g of activated carbon A.
 Part of the activated carbon thus treated was sampled and heated in air for 1 hour at different temperatures of 90°C, 120°C, 150°C and 200°C, respectively.
- 10 (2) Activated carbon having as supported thereon sulfur C₀ through C₂: 10
 90 ml of water was sprayed evenly on 100 g of activated carbon A, and 10.0 g of finely powdered sulfur was sprayed evenly over the wet activated carbon with stirring to coat the particle-surface of the activated carbon with sulfur.
 Part of the activated carbon thus treated was sampled and heated in nitrogen gas for 1/2 hour at
 15 different temperature of 120°C and 250°C, respectively. 15
- (3) Activated carbon having as supported thereon sulfur D:
 90 ml of water was sprayed evenly on 100 g of activated carbon A, and 10.0 g of finely powdered sulfur was sprayed evenly over the wet activated carbon with stirring to coat the particle-surface of the activated carbon with sulfur.
 20 Part of the activated carbon thus treated was sampled and heated in heavy-oil combustion waste 20
 gas at 150°C for 1 hour.
- (4) Activated carbon having as supported thereon sulfur E₀.
 10.0 g of finely powdered sulfur of not more than 50 mesh was dissolved in 120 ml of carbon disulfide, and the solution was sprayed evenly on 100 g of activated carbon A.
 25 Part of the activated carbon thus treated was sampled and heated in nitrogen gas at a 25
 temperature of 80°C for 1/2 hour.
- (5) Activated carbon having as supported thereon sulfur F₀:
 15 g of activated carbon A was packed into a 2.0 cm³ quartz glass tube, which was then heated at 140°C. Air of 140°C containing 1.0 vol.% of H₂S was passed through the packed layer at a linear flow
 30 rate of 15 cm/sec (at 25°C) for 15 hours to have 1.5 g of sulfur supported on the whole activated 30
 carbon.
 The amount of the sulfur supported was found to decrease in the order of the gas-inlet side, middle and gas-outlet side in the packed layer of activated carbon, with the found amounts as sulfur of 14 g/100 g of activated carbon, 10 g/100 g of activated carbon and 6 g/100 g of activated carbon
 35 respectively. The middle layer is referred to as "F₀". 35
 6 ml each of the activated carbon samples A, B₀ through B₃, C₀ through C₂, D, E₀ and F₀ as obtained by the above procedures were packed into a 1.6 cm³ glass column, respectively, and air of 25°C containing 2.0 mg/m³ of mercury vapor was passed through each column at a linear flow rate of 20 cm/sec to conduct the breakthrough adsorption tests for mercury vapor. The results are as shown in
 40 Table 6. 40

TABLE 6

Sulfur-supporting method	Sample No.	Heating temp. °C	Observed conditions of sulfur supported	Removal of mercury vapor at the elapsed time shown below (%)			
				2 hr.	5 hr.	10 hr.	20 hr.
Sulfur is suspended in water, and the suspension is sprayed on activated carbon, which is heated.	B ₀ (Control)	90	Sulfur particles are adhered to external surface	10	4	0	0
	B ₁ (This invention)	120	Sulfur is settled and adhered evenly inside pores.	100	70	50	30
	B ₂ (")	150	"	100	80	63	43
	B ₃ (")	200	"	100	100	65	45
Activated carbon is wetted on the surface with water, then coated with powdered sulfur and heated.	C ₀ (Control)	80	Sulfur particles are adhered to external surface.	13	5	0	0
	C ₁ (This invention)	120	Sulfur is settled and adhered evenly inside pores.	100	65	43	31
	C ₂ (")	250	"	100	78	61	42
	D (This invention)	150	"	100	83	75	60
Sulfur is dissolved in carbon disulfide, and the solution is sprayed on activated carbon, which is heated.	E ₀ (Control)	80	Sulfur is settled and adhered almost evenly inside pores.	70	51	32	21
	F ₀ (Control)	140	The same as above, but the amount of sulfur supported varies among particles of activated carbon.	95	65	48	29
Not treated	A (Control)	—	—	5	0	0	0

EXAMPLE 7

From activated carbon G of 10 to 24 mesh having the BET surface area of 950 m²/g, there was prepared the activated carbon having as supported thereon a mixture of sulfur and other components.

- (1) Activated carbon H having as supported thereon S—KI
- 5 45 ml of an aqueous solution of 1.0 g of KI was evenly sprayed on 100 g of activated carbon G with stirring to allow KI to adhere to the activated carbon. 5 g of sulfur of not more than 20 mesh was poured onto the activated carbon, and stirring was continued for about 10 minutes to coat the surface of the activated carbon with sulfur. The activated carbon as obtained by the above procedure was heated in air at 130°C for 2 hours. 5
- 10 (2) Activated carbon I having as supported thereon S—KI—FeSO₄ 10
- 45 ml of an aqueous solution containing 1.0 g of KI and 2.0 g of FeSO₄ was evenly sprayed on 100 g of activated carbon G with stirring to allow KI and FeSO₄ to adhere to the activated carbon. 3 g of sulfur of not more than 20 mesh was poured onto the activated carbon, and stirring was continued for about 10 minutes to coat the surface of the activated carbon with sulfur. The activated carbon as
- 15 obtained by the above procedure was heated in air at 150°C for 2 hours. 15
- (3) Activated carbon J having as supported thereon S—KBr—Fe(NO₃)₃:
- 48 ml of aqueous ethanol (5 ml of ethanol with the remaining 43 ml of water. Sulfur contained in the state of suspension) containing 1.0 g of KBr, 2.0 g of Fe(NO₃)₃ and 3.0 g of sulfur of not more than 100 mesh was evenly sprayed on 100 g of activated carbon G with stirring, and then heated in an
- 20 atmosphere of nitrogen gas at 115°C for 1 hour, whereby each of the activated carbon samples was found to have sulfur settled and adhered almost uniformly inside pores of the activated carbon. 20
- 25 ml each of the activated carbon samples H, I and J as obtained by the above procedures was packed into a 4 cm⁶ glass tube, and nitrogen gas of 25°C containing 5.0 mg/m³ of mercury vapor was passed through each tube at a linear flow rate of 60 cm/sec to conduct the breakthrough adsorption
- 25 tests for mercury vapor. The results are as shown in Table 7. 25

TABLE 7

Sample No.	Component supported	Percent removal of mercury vapor at the elapsed time shown below			
		1 hr.	5 hrs.	10 hrs.	20 hrs.
H	S—KI	100	95	71	35
I	S—KI—FeSO ₄	100	100	93	65
J	S—KBr—Fe(NO ₃) ₃	100	100	90	52

CLAIMS

1. A process for the removal of mercury vapor from a gas containing the same, which comprises contacting the gas with an adsorbent comprising an activated carbon having as supported thereon (1) one or more components included in one of the three groups shown below and (2) one or more components included in the remaining two groups shown below. 30

Group	Component
I	sulfur
II	sulfate and nitrate of Al, V, Fe, Co, Ni, Cu, Zn or NH ₄
III	oxide of iodine, oxyacid corresponding to the oxide of iodine, salt of said oxyacid, and bromide and iodide of K, Na or NH ₄

2. A process as claimed in Claim 1, wherein the one or more components included in one of the

three groups are sulfur included in Group I, and the one or more components included in the remaining two groups are one or more components included in Group II.

3. A process as claimed in Claim 1, wherein the one or more components included in one of the three groups are sulfur included in Group I, and the one or more components included in the remaining two groups are one or more components included in Group III.

4. A process as claimed in Claim 1, wherein the one or more components included in the other two groups are one or more components included in one of the remaining two components and one or more components included in the other of the remaining two components.

5. A process as claimed in Claim 1, wherein each of the components is supported on the activated carbon so that the amount of sulfur, when used, is in the range of from 10 to 800 mg per g of activated carbon, the total amounts of the components included in Group II, when used, is in the range of from 5 to 500 mg per g of activated carbon, and the total amounts of the components included in Group III, when used, is in the range of from 1 to 500 mg per g of activated carbon.

6. A process as claimed in Claim 1, wherein the activated carbon is one having as supported thereon sulfur and potassium iodide.

7. A process as claimed in Claim 1, wherein the activated carbon is one having as supported thereon sulfur and ferrous sulfate.

8. A process as claimed in Claim 1, wherein the activated carbon is one having as supported thereon sulfur, potassium iodide and ferrous sulfate.

9. An adsorbent for the removal of mercury vapor, which comprises an activated carbon having as supported thereon (1) one or more components included in one of the three groups shown below and (2) one or more components included in the remaining two groups shown below.

Sample	Component
I	sulfur
II	sulfate and nitrate of Al, V, Fe, Co, Ni, Cu, Zn or NH ₄
III	oxide of iodine, oxyacid corresponding to the oxide of iodine, salt of said oxyacid, and bromide and iodide of K, Na. or NH ₄

10. An adsorbent as claimed in Claim 9, wherein the one or more components included in one of the three groups are sulfur included in Group I, and the one or more components included in the remaining two groups are one or more components included in Group II.

11. An adsorbent as claimed in Claim 9, wherein the one or more components included in one of the three groups are sulfur included in Group I, and the one or more components included in the remaining two groups are one or more components included in Group III.

12. An adsorbent as claimed in Claim 9, wherein the one or more components included in the other two groups are one or more components included in one of the remaining two components and one or more components included in the other of the remaining two components.

13. An adsorbent as claimed in Claim 9, wherein each of the components is supported on the activated carbon so that the amount of sulfur, when used, is in the range of from 10 to 800 mg per g of activated carbon, the total amounts of the components included in Group II, when used, is in the range of from 5 to 500 mg per g of activated carbon, and the total amounts of the components included in Group III, when used, is in the range of from 1 to 500 mg per g of activated carbon.

14. An adsorbent as claimed in Claim 9, wherein the activated carbon is one having as supported thereon sulfur and potassium iodide.

15. An adsorbent as claimed in Claim 9, wherein the activated carbon is one having as supported thereon sulfur and ferrous sulfate.

16. An adsorbent as claimed in Claim 9, wherein the activated carbon is one having as supported thereon sulfur, potassium iodide and ferrous sulfate.

17. A process for producing an activated carbon having sulfur as supported thereon, which comprises mixing activated carbon with finely powdered sulfur and heating the resultant mixture at 110 to 350°C for not less than 30 minutes.

18. A process as claimed in Claim 17, wherein the resultant mixture is heated at 110 to 300°C.

19. A process as claimed in Claim 1, substantially as hereinbefore described with reference to any of the Examples.

20. An adsorbent as claimed in Claim 9, substantially as hereinbefore described with reference to any of the Examples.

21. A process as claimed in Claim 17, substantially as hereinbefore described with reference to any of the Examples.

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